Date: 08/08/2005

### IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

1

Applicant:

Ravindra Pandey et al.

U.S. Patent Application No. 10/607,922

For: FLUORINATED PHOTOSENSITIZERS RELATED TO CHLORINS AND BACTERIOCHLORINS FOR PHOTDYNAMIC THERAPY

Filed: June 27, 2003

Examiner: Nwaonicha, Chukwuma

Group Art Unit:

1621

Confirmation No.:

8140

Customer No.: 24041

#### Certificate of Mailing by First Class Mail

I certify that this Correction Letter is being deposited on August 8, 2005 with the U.S. Postal Service as first class mail under 37 C.F.R. §1.8 and is addressed to the Commissioner for Patents, PO Box 1450, Alexandria, VA 22313-1450.

Michael L. Dunn, Reg. No. 25330

### **CORRECTION LETTER**

Mail Stop Amendment Commissioner for Patents PO Box 1450 Alexandria, VA 22313-1450

Honorable Sir:

This calls attention to minor matters on the amendment mailed to the U.S. Patent and Trademark Office on August 4, 2005 (copy of same enclosed in its entirety).

The application number in the header should have been "10/607,922" rather than "10/606,922".

The total claims should have been "19" on the Transmittal Letter instead of "18"...

In the first line of the amendment after "Honorable Sir", "October 20, 2003" should have been "May 4, 2005".

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In view of the foregoing amendments and remarks, it is clear that all claims are in condition for allowance, which action is courteously requested.

Respectfully submitted,

Michael L. Dunn

Registration number 25,330

CUSTOMER NO. 24041

Simpson & Simpson, PLLC

5555 Main Street

Williamsville, NY 14221-5406

Telephone No. 716-626-1564

Dated: August 8, 2005

MLD/mjk

Enc.

OIP	TRANSMIT (General - P	Docket No. RPP174AUS								
In Re Application Of: Chavindra Pandey et al.										
Application No.	Filing Date June 27, 2003	Examiner Chukwuma Nwaonicha	Customer No. 24041	Group Art Unit 1621	Confirmation No. 8140					
Title: FLUORINATED PHOTOSENSITIZERS RELATED TO CHLORINS AND BACTERIOCHLORINS FOR PHOTODYNAMIC THERAPY										
		COMMISSIONER FOR PATI	ENTS:							
Transmitted herew	vith is:				1					
1) Correction Letter 1) Copy of Amendment & Request for Reconsideration dated August 4, 2004 (with copies of transmittal & postcard) 1) Certificate of Mailing by First Class Mail 1) Acknowledgement postcard										
in the above identified application.										
<ul> <li>No additional fee is required.</li> <li>A check in the amount of is attached.</li> <li>The Director is hereby authorized to charge and credit Deposit Account No. 50-0822 as described below.</li> <li>Charge the amount of</li> <li>Credit any overpayment.</li> <li>Charge any additional fee required.</li> <li>Payment by credit card. Form PTO-2038 is attached.</li> <li>WARNING: Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.</li> </ul>										
Mu	Signature		Dated: Aug	ust 8, 2005						
Michael L. Dunn Reg. No. 25330 Simpson & Simpson 5555 Main Street Williamsville, New (716) 626-1564 Pho (716) 626-0366 Fax	York 14221 one		deposited with sufficient postal addressed to the 1450, Alexandria August 8, (Date)	h the United States age as first class the "Commissioner for a, VA 22313-1450" [ 2005-	Correspondence					

Typed or Printed Name of Person Mailing Correspondence

### RECEIVED IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant(s)! Ravindra K. Pandey et al.
U.S. Patent Application No. 10/607,922
Filing Date: June 27, 2003
For: FLUORINATED PHOTOSENSITIZERS RELATED TO CHLORINS AND BACTERIOCHLORINS FOR PHOTODYNAMIC THERAPY

GAU: 1621

Confirmation No.: 8140

Transmitted Herewith is:

(1) Amendment Transmittal Letter (Small Entity)
 (1) Amendment and Request for Reconsideration
 (1) Certificate of Mailing by First Class Mail
 (1) Acknowledgement Postcard

Attorney Docket No.: 1210.RPP174AUS



Customer No. 24041



AMENDMENT TRANSMITTAL LETTER (Small Entity)  Applicant(s): Ravindra K. Pandey et al.						Docket No. RPP174AUS					
1777403											
Application No.	Filing Date	Examiner	Cu	stomer No.	Group Art Unit	Confirmation No.					
10/607,922	June 27,2003	Chukwuma O. Nwaonicha		24041	1621	8140					
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☐ Applicant claims small entity status. See 37 CFR 1.27											
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The fee has been calculated and is transmitted as shown below.											
CLAIMS AS AMENDED											
	CLAIMS REMAINING	HIGHEST#	NUMBER E	XTRA		ADDITIONAL					
	AFTER AMENDMENT	PREV. PAID FOR	CLAIMS PRE	ESENT	RATE	FEE					
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communication or credit any overpayment to Deposit Account No. 50-0822											
Any additional filing fees required under 37 C.F.R. 1.16.											
Any patent application processing fees under 37 CFR 1.17.											
<ul> <li>Payment by credit card. Form PTO-2038 is attached.</li> <li>WARNING: Information on this form may become public. Credit card information should not be</li> </ul>											
included on this form. Provide credit card information and authorization on PTO-2038.											
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Dated: August 4, 2005											
Michael L. Dunn I certify that this correspondence is being deposited with the											
Reg. No. 25330	•		United State	at this corre es Postal Ser	espondence is bein vice with sufficient	g deposited with the postage as first class					
Simpson & Simpson, PLLC mail in an envelope addressed to "Commissioner for Patents, P.O.											
5555 Main Street Box 1450, Alexandria, VA 22313-1450" [37 CFR 1.8(a)] on August 4, 2005											
Williamsville, New York 14221											
(716) 626-1564 Phone (716) 626-0366 Fax											
Muhuel a)											
Signature of Person Mailing Correspondence											

CC:

Michael L. Dunn

Typed or Printed Name of Person Mailing Correspondence



Date: 08/04/2005

### IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant:

Ravindra Pandey et al

U.S. Patent Application No. 10/606,922

For: FLUORINATED

PHOTOSENSITIZERS RELATED TO CHLORINS AND

BACTERIOCHLORINS FOR PHOTDYNAMIC THERAPY

Filed: June 27, 2003

Examiner: Nwaonicha, Chukwuma

Group Art Unit:

1621

Confirmation No.:

8140

Customer No.: 24041

I certify that this Amendment and Request for Reconsideration is being deposited on August 4, 2005 with the U.S. Postal Service as first class mail under 37 C.F.R. §1.8 and is addressed to the Commissioner for Patents, PO Box 1450, Alexandria, VA 22313-1450.

### AMENDMENT AND REQUEST FOR RECONSIDERATION

Mail Stop Amendment Commissioner for Patents PO Box 1450 Alexandria, VA 22313-1450

Honorable Sir:

application as follows:

Responsive to the official action of October 20, 2003, please amend the above identified patent

### In The Claims

Please amend the claims as follows:

Claims 1-4 (cancelled)

Claim 5 (currently amended) A compound of the formula:

or a phamaceutically acceptable derivative thereof, wherein:

 $R_1$  and  $R_2$  are each independently substituted or unsubstituted alkyl, substituted or unsubstituted alkenyl,  $-C(O)R_a$  or  $-COOR_a$  or  $[[-CH(CH_3)(OR) \text{ or } -CH(CH_3)(O(CH_2)_nXR)]]$   $-CH(CH_3)(OR_a)$  or  $-CH(CH_3)(O(CH_2)_nXR_a)$  where  $R_a$  is hydrogen, substituted or unsubstituted alkyl, substituted or unsubstituted or unsubstituted alkynyl, or substituted or unsubstituted alkynyl, or substituted or unsubstituted cycloalkyl where  $R_2$  may be  $CH-CH_2$ ,  $CH(OR_{20})CH_3$ , C(O)Me,  $C(-NR_{21})CH_3$  or  $CH(NHR_{21})CH_3$ ; where  $R_2$  may be  $-CH=CH_2$ ,  $-CH(OR_{20})CH_3$ , -C(O)Me,  $-C(=NR_{21})CH_3$  or  $-CH(NHR_{21})CH_3$ 

where X is an aryl or heteroaryl group:

n is an integer of 0 to 6;

-----R and R

where  $R_{20}$  is methyl, butyl, heptyl, docecyl or 3,5-bis(trifluoromethyl)-benzyl; and  $R_{21}$  is 3,5,-bis(trifluoromethyl)benzyl;

 $R_{1a}$  and  $R_{2a}$  are each independently hydrogen or substituted or unsubstituted alkyl, or together form a covalent bond;

R<sub>3</sub> and R<sub>4</sub> are each independently hydrogen or substituted or unsubstituted alkyl;

 $R_{3a}$  and  $R_{4a}$  are each independently hydrogen or substituted or unsubstituted alkyl, or together form a covalent bond;

R<sub>5</sub> is hydrogen or substituted or unsubstituted alkyl;

 $R_6$  and  $R_{6a}$  are each independently hydrogen or substituted or unsubstituted alkyl, or together form =0:

 $R_7$  is a covalent bond, alkylene, azaalkyl, or azaaraalkyl or =NR<sub>20</sub> where R<sub>20</sub> is 3,5-bis(tri-fluoromethyl)benzyl or -CH<sub>2</sub>X-R<sup>1</sup> or -YR<sup>1</sup> where Y is an aryl or heteroaryl group;

 $R_8$  and  $R_{8a}$  are each independently hydrogen or substituted or unsubstituted alkyl or together form =0;

R<sub>9</sub> and R<sub>10</sub> are each independently hydrogen, or substituted or unsubstituted alkyl and R<sub>9</sub> may be -CH<sub>2</sub>CH<sub>2</sub>COOR<sup>2</sup> where R<sup>2</sup> is an alkyl group that may optionally substituted with one or more fluorine atoms;

each of R<sub>1</sub>-R<sub>10</sub>, when substituted, is substituted with one or more substituents each independently selected from Q, where Q is alkyl, haloalkyl, halo, pseudohalo, or -COOR<sub>b</sub> where R<sub>b</sub> is hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, er aryl, heteroaryl, araalkyl, or OR<sub>c</sub> where R<sub>c</sub> is hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, or aryl or CONR<sub>d</sub>R<sub>e</sub> where R<sub>d</sub> and R<sub>e</sub> are each independently hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, or aryl, or NR<sub>f</sub>R<sub>g</sub> where R<sub>f</sub> and R<sub>g</sub> are each independently hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, or aryl, or aryl, or =NR<sub>h</sub> where R<sub>h</sub> is hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, or is an amino acid residue;

each Q is independently unsubstituted or is substituted with one or more substituents each independently selected from  $Q_1$ , where  $Q_1$  is alkyl, haloalkyl, halo, pseudohalo, or -COOR<sub>b</sub> where  $R_b$  is hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, er aryl, heteroaryl, araalkyl, or  $OR_c$  where  $R_c$  is hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, or aryl or  $CONR_dR_c$  where  $R_d$  and  $R_c$  are each independently hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, or aryl, or  $NR_fR_g$  where  $R_f$  and  $R_g$  are each independently hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, or aryl, or aryl, or  $PR_f$  where  $PR_f$  is hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, or aryl, or is an amino acid residue.

with the proviso that the compound contains at least one fluorine atom in at least one

3,5,-bis(trifluoromethyl)benzyl group or in at least one R, R<sup>1</sup>, or R<sup>2</sup> group.

Claims 6-7 (cancelled)

Claim 8 (previously presented). The compound of claim 5 wherein:

R<sub>1</sub> is methyl;

R<sub>1a</sub> and R<sub>2a</sub> together form a covalent bond;

R<sub>3</sub> is methyl;

R<sub>4</sub> is ethyl;

 $R_{3a}$  and  $R_{4a}$  are each independently hydrogen, or together form a covalent bond;

R₅ is methyl;

R<sub>9</sub> is CH<sub>2</sub>CH<sub>2</sub>COOH or CH<sub>2</sub>CH<sub>2</sub>COOMe;

R<sub>10</sub> is methyl.

Claim 9 (previously presented) The compound of claim 5, wherein:

 $R_2$  is CH=CH<sub>2</sub>, CH(OR<sub>20</sub>)CH<sub>3</sub>, C(O)Me, C(=NR<sub>21</sub>)CH<sub>3</sub> or CH(NHR<sub>21</sub>)CH<sub>3</sub>;

where  $R_{20}$  is methyl, butyl, heptyl, dodecyl or 3,5-bis(trifluoromethyl)-benzyl; and

R<sub>21</sub> is 3,5-bis(trifluoromethyl)benzyl.

Date: 08/04/2005

Claim 10 (cancelled)

Claim 11 (previously presented) The compound of claim 5 having the formula:

or a pharmaceutically acceptable derivative thereof.

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### Claim 12 (previously presented) The compound of claim 5 having the formula:

or a pharmaceutically acceptable derivative thereof, wherein:

R is methyl, butyl, heptyl or dodecyl.

## Claim 13 (previously presented) The compound of claim 5 having the formula:

or a pharmaceutically acceptable derivative thereof, wherein:

R is methyl, butyl, heptyl or dodecyl.

# Claim 14 (previously presented) The compound of claim 5 having the formula:

or a pharmaceutically acceptable derivative thereof.

# Claim 15 (previously presented) The compound of claim 5 having the formula:

or a pharmaceutically acceptable derivative thereof, wherein:

R is methyl, butyl, heptyl or dodecyl.

## Claim 16 (previously presented) The compound of claim 5 having the formula:

or a pharmaceutically acceptable derivative thereof.

Claim 17 (previously presented) The compound of claim 5 having the formula:

or a pharmaceutically acceptable derivative thereof.

### Claim 18 (previously presented) The compound of claim 5 having the formula:

or a pharmaceutically acceptable derivative thereof, wherein:

X is an aryl or heteroaryl group;

R and  $R^1$  are each independently alkyl, aryl, or heteroaryl groups having 1 – 20 carbon atoms, wherein at least one of R and  $R^1$  is substituted with at least one fluorine atom; and

R<sup>2</sup> is an alkyl group, optionally substituted with one or more fluorine atoms.

Claim 19 (previously presented) The compound of claim 5 having the formula:

or a pharmaceutically acceptable derivative thereof, wherein:

X is an aryl or heteroaryl group;

n is an integer from 0 to 6;

R and  $R^1$  are each independently alkyl, aryl, or heteroaryl groups having 1 – 20 carbon atoms, wherein at least one of R and  $R^1$  is substituted with at least one fluorine atom; and

 $\ensuremath{\mathbb{R}}^2$  is an alkyl group, optionally substituted with one or more fluorine atoms.

Claim 20 (previously presented) The compound of claim 5 having the formula

or a pharmaceutically acceptable derivative thereof, wherein:

X is an aryl or heteroaryl group;

R and  $R^1$  are each independently alkyl, aryl, or heteroaryl groups having 1-20 carbon atoms, wherein at least one of R and  $R^1$  is substituted with at least one fluorine atom; and

R<sup>2</sup> is an alkyl group, optionally substituted with one or more fluorine atoms.

Claim 21 (previously presented) The compound of claim 5 having the formula:

or a pharmaceutically acceptable derivative thereof, wherein:

X is an aryl or heteroaryl group;

n is an integer from 0 to 6;

R and  $R^1$  are each independently alkyl, aryl, or heteroaryl groups having 1 – 20 carbon atoms, wherein at least one of R and  $R^1$  is substituted with at least one fluorine atom; and

R<sup>2</sup> is an alkyl group, optionally substituted with one or more fluorine atoms.

Date: 08/04/2005

# Claim 22 (previously presented) The compound of claim 5 having the formula:

or a pharmaceutically acceptable derivative thereof, wherein:

X and Y are each independently an aryl or heteroaryl group; n is an integer from 0 to 6;

R and  $R^1$  are each independently alkyl, aryl, or heteroaryl groups having 1 – 20 carbon atoms, wherein at least one of R and  $R^1$  is substituted with at least one fluorine atom; and

R<sup>2</sup> is an alkyl group, optionally substituted with one or more fluorine atoms.

Claim 23 (previously presented)

A pharmaceutical composition, comprising a compound of claim 1 or a pharmaceutically acceptable derivative thereof in a pharmaceutically acceptable carrier.

Claims 24-121 (cancelled)

Date: 08/04/2005

Claim 122 (previously presented) The compound of claim 17 or a pharmaceutically acceptable derivative thereof when used for the detection or treatment or both of hyperproliferative tissue.

Claim 123 (previously presented) The compound of claim 18 or a pharmaceutically acceptable derivative thereof when used for the detection or treatment or both of hyperproliferative tissue.

Claim 124 (previously presented) The compound of claim 19 or a pharmaceutically acceptable derivative thereof when used for the detection or treatment or both of hyperproliferative tissue.

#### Remarks

The Examiner in the official action referred a couple of times to a response filed by the Applicants on December 13, 2004. The Applicants filed no such amendment on that at but rather filed an amendment on February 7, 2005. It is unclear what the Examiner is referring to.

The Examiner has rejected claim 8 as being anticipated by the Li et al reference under 35 U.S.C. 102. The Examiner is in error.

The amendment of February 7, 2005 amended claim 5 so that it requires at least one fluorine atom in at least one 3,5,-bis(trifluoromethyl)benzyl group or in at least one R, R<sup>1</sup>, or R<sup>2</sup> group. All remaining claims depend from claim 5 and thus also require at least one fluorine atom in at least one 3,5,-bis(trifluoromethyl)benzyl group or in at least one R, R<sup>1</sup>, or R<sup>2</sup> group. Claim 8 thus requires at least one fluorine atom in at least one 3,5,-bis(trifluoromethyl)benzyl group or in at least one R, R<sup>1</sup>, or R<sup>2</sup> group. Li et al discloses or suggests no such compound.

The rejection should be withdrawn.

Claims 5, 8, 9 and 10 have been rejected under 35 U.S.C. 103 as being unpatentable over Pandey et al., U.S. 5,952,366) in view of Li et al.

This rejection is improper and should be withdrawn.

As discussed above, All remaining claims require at least one fluorine atom in at least one 3,5,-bis(trifluoromethyl)benzyl group or in at least one R, R<sup>1</sup>, or R<sup>2</sup> group. Pandey et al. disclose or suggest no such compound. No trifluoromethyl compound of any kind is

Date: 08/04/2005

suggested by Pandey et al. and likewise no trifluoromethyl compound of any kind is suggested by Li et al. It is thus clear that this combination of references cannot and does not suggest any trifluoromethyl compound and certainly not the (trifluoromethyl)benzyl compounds

presently claimed.

The Examiner has rejected claim 5 as being indefinite under 35 U.S.C. 112 because R

and R' are not defined. R and R' and have been deleted by amendment.

The Examiner has rejected claim 5 as being indefinite under 35 U.S.C. 112 because the nitrogen in the general formula has two bonds instead of three. This is clearly a typographical type error. The nitrogens in the a and c rings clearly have an attached hydrogen as is well known with respect to unsaturated a and c rings of tetrapyrol type compounds. The claim has been amended to remove the objection.

Claim 10 has been cancelled.

In view of the foregoing amendments and remarks, it is clear that all claims are in condition for allowance, which action is courteously requested.

Respectfully submitted

Michael L. Dunn

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